## CLAIM AMENDMENTS

## 1. (canceled)

- 2. (currently amended) A method of making a vascular prosthesis or tissue web of biocompatible polymer, especially of polyurethane, polyamide, polysulfone, polyester, isotactic polypropylene, polynitrile [[and/]] or polyvinylchloride, (currently amended) mixtures thereof [[and/]] or their copolymers, with a microporous finely fibular structure, characterized by a definitive stretching (extension) with a degree of extension between 30% and 150% preferably between 60 and (currently amended) 125%, and subsequent relaxation.
- 3. (currently amended) The method according to claim 2

  characterized in that the wherein a pore size of the vascular

  prosthesis or of the tissue patch before the stretching is less

  than [[the]] an extended dimension expected prior to stretching and

  beyond which the vascular prosthesis or tissue patch does not

  retract.
- 4. (currently amended) The method according to claim 2

  characterized in that wherein the stretching is [[an]] a uniaxial

  or biaxial stretching.

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- 5. (currently amended) The method according to claim 27
  characterized in that wherein the vascular prosthesis or the tissue
  patch prior (previously presented) to the stretching is soaked in a
  water soluble polyphysiological substance, preferably
  polyvinylalcohol (PVA), polyvinylpyrrolidone or gelatine (collagen)
  [[which]] that is completely or partially drawn into the vascular
  prosthesis or the tissue patch, preferably on [[the]] an outer
  side thereof.
- 6. (currently amended) The method according to claim 27
  characterized in that wherein the vascular prosthesis is tubular
  and for stretching a requisite pressure is applied from the
  interior with a gaseous medium, preferably air or N2, or with a
  liquid medium (previously presented).
  - 7. (currently amended) The method according to claim 6 characterized in that wherein to avoid leakage, a yieldable preferably elastic auxiliary body is introduced into the vascular prosthesis to be stretched and is thereafter pressurized with a pressure applying medium.
    - 8. (currently amended) The method according to claim 57 characterized in that wherein the stretching is carried out with an auxiliary body capable of mechanical size adjustment upon which the

- tissue patch is previously clamped or which is introduced into the
- 5 tubular prosthesis.
- 9. (currently amended) The method according to claim 57
- 2 characterized in that wherein for widening a tubular vascular
- prosthesis, a drawing mandrel is used.
- 10. (currently amended) The method according to claim 27
- 2 characterized in that wherein to produce the vascular prosthesis or
- 3 the tissue patch at least one aliphatic and/or at least one
- 4 cycloaliphatic diisocyanate is reacted with a polyarbonate,
- 5 polyester, polyether, polysiloxane, or polysulfone macrodiol of the
- 6 polycarbonate type or of the polyester, (previously presented)
- 7 polyether, polysiloxane or polysulfone type with an average
- s molecular weight of 500 to 6000, whereby the ratio of NCO terminal
- groups of the prepolymer to OH groups of the chain lengthening
- agent is 1.01:1 to 1.05:1 and the polymer obtained, optionally
- aftertreatment with a reagent for deactivating NCO groups which may
- still be present, is subjected to a molecular weight fractionation
- in which the low molecular weight polyurethane fraction making up
- 10% to 50% by weight of the polymer (previously presented) is
- separated off and discarded and the remaining high molecular weight
- fractionation is recovered as the biocompatible polyurethane with
- improved properties.

- 1 11. (new) The method according to claim 2 wherein the
- degree of extension is 60% to 125%.
- 12. (new) The method according to claim 2 wherein the
- prosthesis or web is relaxed by 3% to 5%.